

# Complete mitochondrial genome of *Conus lischkeanus* Weinkauff, 1875 (Neogastropoda, Conidae) and phylogenetic implications of the evolutionary diversification of dietary types of *Conus* species

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## Abstract

The family Conidae, commonly known as cone snails, is one of the most intriguing gastropod groups owing to their diverse array of feeding behaviors (diets) and toxin peptides (conotoxins). *Conus lischkeanus* Weinkauff, 1875 is a worm-hunting species widely distributed from Africa to the Northwest Pacific. In this study, we report the mitochondrial genome sequence of *C. lischkeanus* and inferred its phylogenetic relationship with other *Conus* species. Its mitochondrial genome is a circular DNA molecule (16,120 bp in size) composed of 37 genes: 13 protein-coding genes (PCGs), 22 transfer RNA genes, and two ribosomal RNA genes. Phylogenetic analyses of concatenated nucleotide sequences of 13 PCGs and two ribosomal RNA genes showed that *C. lischkeanus* belongs to the subgenus *Lividoconus* group, which is grouped with species of the subgenus *Virgiconus*, and a member of the largest assemblage of worm-hunting (vermivorous) species at the most basal position in this group. Mitochondrial genome phylogeny supports the previous hypothesis that the ancestral diet of cone snails was worm-hunting, and that other dietary types (molluscivorous or piscivorous) have secondarily evolved multiple times from different origins. This new, complete mitochondrial genome information provides valuable insights into the mitochondrial genome diversity and molecular phylogeny of *Conus* species.



**Keywords**

Cone snail, dietary type evolution, *Lividoconus*

**Introduction**

The genus *Conus* Linnaeus, 1758 is a well-known predatory gastropod group that produces venomous peptides, called conotoxins, to capture prey and defend against predators (Dutertre et al. 2004; Prashanth et al. 2016; Kohn 2019). There are more than 750 *Conus* species reported worldwide (WoRMS 2021), which are widely distributed in tropical and subtropical ocean areas in various environments ranging from deep water to the intertidal zone (Kohn 1959). With the notable exception of a few conid species that prey on more than one dietary type (e.g., *Californiconus californicus* (Reeve, 1844) and *Conus bullatus* Linnaeus, 1758), most species in this genus show a very narrow range of prey, feeding on worms, mollusks, and fishes, and they are grouped into three specialized dietary types according to their prey types: vermivorous (worm-hunting), molluscivorous (mollusk-hunting), and piscivorous (fish-hunting) (Duda Jr et al. 2001; Olivera et al. 2014; Robinson et al. 2014; Himaya et al. 2015; Gao et al. 2018). Among these diverse dietary types, the worm-hunting diet is the most common, accounting for more than 70% of the species, and it is widely considered the most ancestral; other dietary types are regarded to have undergone secondary evolution (Duda Jr et al. 2001; Puillandre et al. 2014; Gao et al. 2018; Abalde et al. 2019). The evolutionary origin and diversification of their dietary specification can be better understood based on well-reconstructed phylogenetic relationships among *Conus* species of different diet types.

The implementation of new sequencing technologies (e.g., next-generation sequencing; NGS) and various bioinformatics tools has allowed mitochondrial genome sequencing to be markedly easier, cost-effective, and widely used for studying phylogeny in various metazoan groups, including *Conus* species (Abalde et al. 2017; Uribe et al. 2017, 2018). As of January 2022, complete and partial mitochondrial genome sequences of 60 *Conus* species have been reported in GenBank, most of which are tropical and subtropical species, and diverse species in other oceanic regions are relatively underrepresented. To elucidate the phylogenetic relationships and evolution of dietary specialization within the genus, phylogenetic analysis using additional mitochondrial genome information sampled from various regional species is needed. To date, only partial mitochondrial gene sequences (12S, 16S, and *cox1*) of *Conus lischkeanus* are currently available on GenBank, with no complete mitochondrial genome information for this species. *Conus lischkeanus* Weinkauff, 1875 is a vermivorous species reported from East Africa to the western Pacific (Röckel et al. 1995). In this study, we determine the complete mitochondrial genome of *C. lischkeanus* for the first time and perform a phylogenetic analysis of 13 protein-coding genes and two ribosomal RNA (rRNA) gene sequences of 39 *Conus* species with different dietary types, including *C. lischkeanus*.



## Materials and methods

### Sample collection and DNA extraction

*Conus lischkeanus* specimen was collected from Moonseom, Jeju Island, Korea, preserved in 95% ethanol solution, and deposited in the Marine Mollusk Resource Bank of Korea (MMRBK; voucher specimen no. MMRBK6746) in Seoul, Korea. The specimen was morphologically identified based on shell characters, which include a conical last whorl covered with yellow-brown periostracum and an angular shoulder. Total genomic DNA was extracted from the foot tissue using an E.Z.N.A. mollusc DNA kit (Omega Bio-tek, Norcross, GA, USA) following the manufacturer's instructions.

### NGS and mitochondrial genome assembly and annotation

Whole-genome sequencing libraries were prepared using the MGIEasy DNA library prep kit (BGI, Shenzhen, China) according to the manufacturer's instructions and quantified using the QuantiFluor ssDNA System (Promega Corporation, Madison, WI, USA). Sequencing was conducted on the MGISEQ-2000 system with 150 base-pair reads. A total of 48,608,637 raw reads were obtained, and adapter-trimmed using a skewer program (Jiang et al. 2014) with a mean quality threshold of 20. The mitochondrial genome was assembled from trimmed reads using MITObim v. 1.9.1 (Hahn et al. 2013). Mitochondrial gene annotation was performed using MITOS webserver (Bernt et al. 2013) and confirmed through sequence comparison with mitochondrial genomes of other *Conus* species previously reported (Bandyopadhyay et al. 2008; Cunha et al. 2009; Brauer et al. 2012; Barghi et al. 2016; Chen et al. 2016a, 2016b, 2016c; Gao et al. 2018; Uribe et al. 2018) using Geneious v. 9.1.8 (Kearse et al. 2012). The nucleotide composition, amino acid composition, and relative synonymous codon usage (RSCU) were analyzed using the MEGA X program (Kumar et al. 2018). Nucleotide composition skew was calculated using the following formula:  $AT\text{-skew} = [A - T] / [A + T]$  and  $GC\text{-skew} = [G - C] / [G + C]$  (Perna and Kocher 1995).

### Phylogenetic analysis

To determine the relationship between *C. lischkeanus* and other *Conus* species, phylogenetic analyses were performed for the nucleotide sequences of 13 protein-coding genes (PCGs) and two rRNA genes from 39 complete or nearly complete mitochondrial genomes of the family Conidae (Table 1). *Tomopleura* sp., belonging to the family Borsoniidae, was also included as an outgroup in the analysis. A concatenated nucleotide sequence dataset (13,870 bp long) of the 13 PCGs and two rRNA genes was prepared for phylogenetic analysis. The best substitution model for each gene was estimated using jModelTest v. 2.1.10 (Darriba et al. 2012) with the Akaike information criterion (AIC) for the nucleotide dataset. Phylogenetic analy-



**Table 1.** Complete mitochondrial genomes used for phylogenetic analysis in this study.

| Family      | Species                            | Diet           | GenBank  | Reference                 |
|-------------|------------------------------------|----------------|----------|---------------------------|
| Conidae     | <i>Conus victoriae</i>             | Molluscivorous | —        | Abalde et al. 2019        |
|             | <i>Conus gloriamaris</i>           | Molluscivorous | KU996360 | —                         |
|             | <i>Conus textile</i>               | Molluscivorous | DQ862058 | Bandyopadhyay et al. 2008 |
|             | <i>Conus episcopatus</i>           | Molluscivorous | —        | Abalde et al. 2019        |
|             | <i>Conus marmoreus</i>             | Molluscivorous | —        | Abalde et al. 2019        |
|             | <i>Conus nobilis</i>               | Molluscivorous | KX263253 | Uribe et al. 2017         |
|             | <i>Conus ermineus</i>              | Piscivorous    | KY864977 | Abalde et al. 2017        |
|             | <i>Conus tulipa</i>                | Piscivorous    | KR006970 | Chen et al. 2016a         |
|             | <i>Conus consors</i>               | Piscivorous    | KF887950 | Brauer et al. 2012        |
|             | <i>Conus striatus</i>              | Piscivorous    | KX156937 | Chen et al. 2016b         |
|             | <i>Conus betulinus</i>             | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus sponsalis</i>             | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus arenatus</i>              | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus goudeyi</i>               | Vermivorous    | KY864975 | Abalde et al. 2019        |
|             | <i>Conus ebraeus</i>               | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus coronatus</i>             | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus miliaris</i>              | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus pseudonivifer</i>         | Vermivorous    | KY864969 | Abalde et al. 2017        |
|             | <i>Conus venulatus</i>             | Vermivorous    | KX263250 | Uribe et al. 2017         |
|             | <i>Conus ateralbus</i>             | Vermivorous    | KY864970 | Abalde et al. 2017        |
|             | <i>Conus byssinus</i>              | Vermivorous    | KY864973 | Abalde et al. 2017        |
|             | <i>Conus pulcher</i>               | Vermivorous    | KY864972 | Abalde et al. 2017        |
|             | <i>Conus genuanus</i>              | Vermivorous    | KY864974 | Abalde et al. 2019        |
|             | <i>Conus hybridus</i>              | Vermivorous    | KX263252 | Uribe et al. 2017         |
|             | <i>Conus guanche</i>               | Vermivorous    | KY801847 | Abalde et al. 2017        |
|             | <i>Conus ventricosus</i>           | Vermivorous    | KX263251 | Uribe et al. 2017         |
|             | <i>Conus miruchae</i>              | Vermivorous    | KY864971 | Abalde et al. 2017        |
|             | <i>Conus borgei</i>                | Vermivorous    | EU827198 | Cunha et al. 2009         |
|             | <i>Conus infinitus</i>             | Vermivorous    | KY864967 | Abalde et al. 2017        |
|             | <i>Conus spurius</i>               | Vermivorous    | KY864976 | Abalde et al. 2019        |
|             | <i>Conus virgo</i>                 | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus quercinus</i>             | Vermivorous    | KY609509 | Gao et al. 2018           |
|             | <i>Conus lischkeanus</i>           | Vermivorous    | OL632021 | This study                |
|             | <i>Conus lividus</i>               | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus tabidus</i>               | Vermivorous    | KY864968 | Abalde et al. 2019        |
|             | <i>Conus lenavati</i>              | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus tribblei</i>              | Vermivorous    | KT199301 | Barghi et al. 2016        |
|             | <i>Conus imperialis</i>            | Vermivorous    | —        | Abalde et al. 2019        |
|             | <i>Conus capitaneus</i>            | Vermivorous    | KX155573 | Chen et al. 2016c         |
|             | <i>Conasprella wakayamaensis</i>   | Vermivorous    | KX263254 | Uribe et al. 2017         |
|             | <i>Californiconus californicus</i> | All            | KX263249 | Uribe et al. 2017         |
|             | <i>Profundiconus teramachii</i>    | Vermivorous    | KX263256 | Uribe et al. 2017         |
| Borsoniidae | <i>Tomopleura</i> sp.              | —              | KX263259 | Uribe et al. 2017         |

ses were conducted using maximum likelihood (ML) and Bayesian inference (BI) methods. ML analysis was performed using RAxML v. 8.2.9 (Stamatakis 2014) with a heuristic search and 10,000 bootstrap replicates. The BI tree was generated using the Markov chain Monte Carlo method, with two independent runs of  $1 \times 10^6$  generations with four chains, sampling every 100 generations and discarding the first 25% generations as burn-in. Both ML and BI programs were conducted using the CIPRES portal (Miller et al. 2010).



Results and discussion

Mitochondrial genome organization and nucleotide composition

*Conus lischkeanus* is widely distributed from East Africa to the western Pacific (Röckel et al. 1995), extending to Taiwan, Japan, and Korea (Jeju Island). This species shows a wide range of shell morph and color variations, depending on geographic origin, which were previously classified as a few separate subspecies (Coomans and Filmer 1985) but are now treated as local variations of *C. lischkeanus* (Röckel et al. 1995). In this study, we determine the complete mitochondrial genome of *C. lischkeanus* and compare it with other cone snail species to infer the evolutionary diversification of different dietary types. The complete mitochondrial genome of *C. lischkeanus* (GenBank accession number: OL632021) is 16,120 bp in size, encoding 13 PCGs, 22 tRNA genes, two rRNA genes, and one control region (Fig. 1, Table 2). The overall nucleotide base composition is 29% A, 37.1% T, 17.6% G, and 16.3% C (Table 3). All 13 PCGs, 14 tRNAs, and two rRNA genes are encoded on the heavy strand, whereas eight tRNA genes (*trnT*, *trnM*, *trnY*, *trnC*, *trnW*, *trnQ*, *trnG*, and *trnE*) are encoded on the light strand. The gene order is identical to that of other cone snail species, suggesting that the mitochondrial

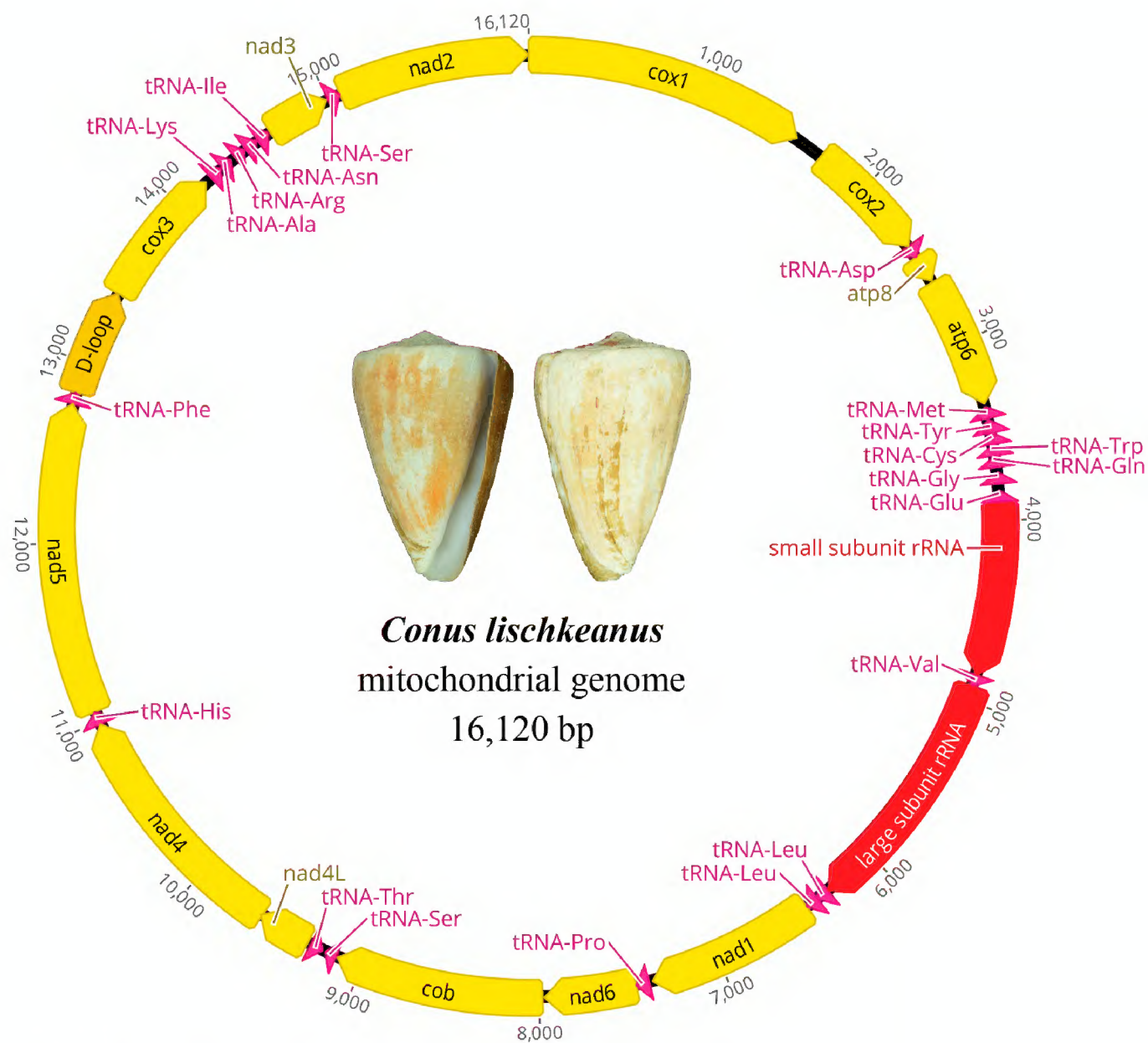


Figure 1. Mitochondrial genome structure of *Conus lischkeanus*.

**Table 2.** Gene regions in the mitochondrial genome of *Conus lischkeanus*.

| Gene                               | Start  | Stop   | Strand<br>direction | Length<br>(bp) | Codon<br>(start) | Codon<br>(stop) | Overlapping<br>regions | Intergenic<br>spacers |
|------------------------------------|--------|--------|---------------------|----------------|------------------|-----------------|------------------------|-----------------------|
| <i>cox1</i>                        | 1      | 1,548  | H                   | 1,548          | ATG              | TAA             | 1                      | 166                   |
| <i>cox2</i>                        | 1,715  | 2,401  | H                   | 687            | ATG              | TAA             | —                      | —                     |
| tRNA-Asp ( <i>trnD</i> ) (gtc)     | 2,402  | 2,468  | H                   | 67             |                  |                 | —                      | —                     |
| <i>atp8</i>                        | 2,469  | 2,630  | H                   | 162            | ATG              | TAA             | —                      | 6                     |
| <i>atp6</i>                        | 2,637  | 3,359  | H                   | 723            | ATG              | TAA             | —                      | 11                    |
| tRNA-Met ( <i>trnM</i> ) (cat)     | 3,371  | 3,438  | L                   | 68             |                  |                 | —                      | 12                    |
| tRNA-Tyr ( <i>trnY</i> ) (gta)     | 3,451  | 3,516  | L                   | 66             |                  |                 | —                      | 1                     |
| tRNA-Cys ( <i>trnC</i> ) (gca)     | 3,518  | 3,582  | L                   | 65             |                  |                 | —                      | —                     |
| tRNA-Trp ( <i>trnW</i> ) (tca)     | 3,583  | 3,648  | L                   | 66             |                  |                 | —                      | —                     |
| tRNA-Gln ( <i>trnQ</i> ) (ttg)     | 3,646  | 3,711  | L                   | 66             |                  |                 | 3                      | 24                    |
| tRNA-Gly ( <i>trnG</i> ) (tcc)     | 3,736  | 3,802  | L                   | 67             |                  |                 | —                      | 35                    |
| tRNA-Glu ( <i>trnE</i> ) (ttc)     | 3,838  | 3,902  | L                   | 65             |                  |                 | —                      | —                     |
| small subunit rRNA ( <i>rrnS</i> ) | 3,903  | 4,854  | H                   | 952            |                  |                 | —                      | —                     |
| tRNA-Val ( <i>trnV</i> ) (tac)     | 4,855  | 4,921  | H                   | 67             |                  |                 | —                      | —                     |
| large subunit rRNA ( <i>rrnL</i> ) | 4,922  | 6,296  | H                   | 1,375          |                  |                 | —                      | —                     |
| tRNA-Leu1 ( <i>trnL1</i> ) (tag)   | 6,297  | 6,366  | H                   | 70             |                  |                 | —                      | 6                     |
| tRNA-Leu2 ( <i>trnL2</i> ) (taa)   | 6,373  | 6,441  | H                   | 69             |                  |                 | —                      | —                     |
| <i>nad1</i>                        | 6,442  | 7,383  | H                   | 942            | ATG              | TAG             | —                      | 16                    |
| tRNA-Pro ( <i>trnP</i> ) (tgg)     | 7,400  | 7,468  | H                   | 69             |                  |                 | —                      | —                     |
| <i>nad6</i>                        | 7,469  | 7,975  | H                   | 507            | ATG              | TAA             | —                      | 13                    |
| <i>cob</i>                         | 7,989  | 9,128  | H                   | 1,140          | ATG              | TAA             | —                      | 11                    |
| tRNA-Ser2 ( <i>trnS2</i> ) (tga)   | 9,140  | 9,204  | H                   | 65             |                  |                 | —                      | 16                    |
| tRNA-Thr ( <i>trnT</i> ) (tgt)     | 9,221  | 9,289  | L                   | 69             |                  |                 | —                      | 22                    |
| <i>nad4L</i>                       | 9,312  | 9,608  | H                   | 297            | ATG              | TAG             | —                      | —                     |
| <i>nad4</i>                        | 9,602  | 10,984 | H                   | 1,383          | ATG              | TAG             | 7                      | —                     |
| tRNA-His ( <i>trnH</i> ) (gtg)     | 10,984 | 11,049 | H                   | 66             |                  |                 | 1                      | —                     |
| <i>nad5</i>                        | 11,050 | 12,765 | H                   | 1,716          | ATG              | TAA             | —                      | —                     |
| tRNA-Phe ( <i>trnF</i> ) (gaa)     | 12,765 | 12,829 | H                   | 65             |                  |                 | 1                      | —                     |
| D-loop                             | 12,830 | 13,415 | H                   | 586            |                  |                 | —                      | —                     |
| <i>cox3</i>                        | 13,416 | 14,195 | H                   | 780            | ATG              | TAA             | —                      | 34                    |
| tRNA-Lys ( <i>trnK</i> ) (ttt)     | 14,230 | 14,298 | H                   | 69             |                  |                 | —                      | 9                     |
| tRNA-Ala ( <i>trnA</i> ) (tgc)     | 14,308 | 14,374 | H                   | 67             |                  |                 | —                      | 22                    |
| tRNA-Arg ( <i>trnR</i> ) (tcg)     | 14,397 | 14,465 | H                   | 69             |                  |                 | —                      | 11                    |
| tRNA-Asn ( <i>trnN</i> ) (gtt)     | 14,477 | 14,545 | H                   | 69             |                  |                 | —                      | 12                    |
| tRNA-Ile ( <i>trnI</i> ) (gat)     | 14,558 | 14,626 | H                   | 69             |                  |                 | —                      | 5                     |
| <i>nad3</i>                        | 14,632 | 14,985 | H                   | 354            | ATG              | TAA             | —                      | 15                    |
| tRNA-Ser1 ( <i>trnS1</i> ) (gct)   | 15,001 | 15,068 | H                   | 68             |                  |                 | —                      | —                     |
| <i>nad2</i>                        | 15,069 | 1      | H                   | 1,053          | ATG              | TAA             | —                      | —                     |

**Table 3.** Nucleotide composition of the mitochondrial genome of *Conus lischkeanus*.

| Nucleotide sequence         | Length (bp) | A (%) | C (%) | G (%) | T (%) | A+T (%) | G+C (%) |
|-----------------------------|-------------|-------|-------|-------|-------|---------|---------|
| Entire sequence             | 16,120      | 29.0  | 16.3  | 17.6  | 37.1  | 66.1    | 33.9    |
| Protein coding sequence     | 11,292      | 26.3  | 17.0  | 17.5  | 39.2  | 65.5    | 34.5    |
| Codon position*             |             |       |       |       |       |         |         |
| 1 <sup>st</sup>             | 3,751       | 26.9  | 17.2  | 24.7  | 31.2  | 58.1    | 41.9    |
| 2 <sup>nd</sup>             | 3,751       | 18.3  | 20.9  | 16.6  | 44.2  | 62.5    | 37.5    |
| 3 <sup>rd</sup>             | 3,751       | 33.4  | 13.1  | 11.4  | 42.1  | 75.5    | 24.5    |
| Ribosomal RNA gene sequence | 2,327       | 35.5  | 14.4  | 18.4  | 31.6  | 67.2    | 32.8    |
| Transfer RNA gene sequence  | 1,481       | 34.0  | 16.2  | 17.7  | 32.1  | 66.1    | 33.9    |
| D-loop region sequence      | 586         | 31.1  | 15.4  | 17.6  | 35.8  | 67.1    | 32.9    |

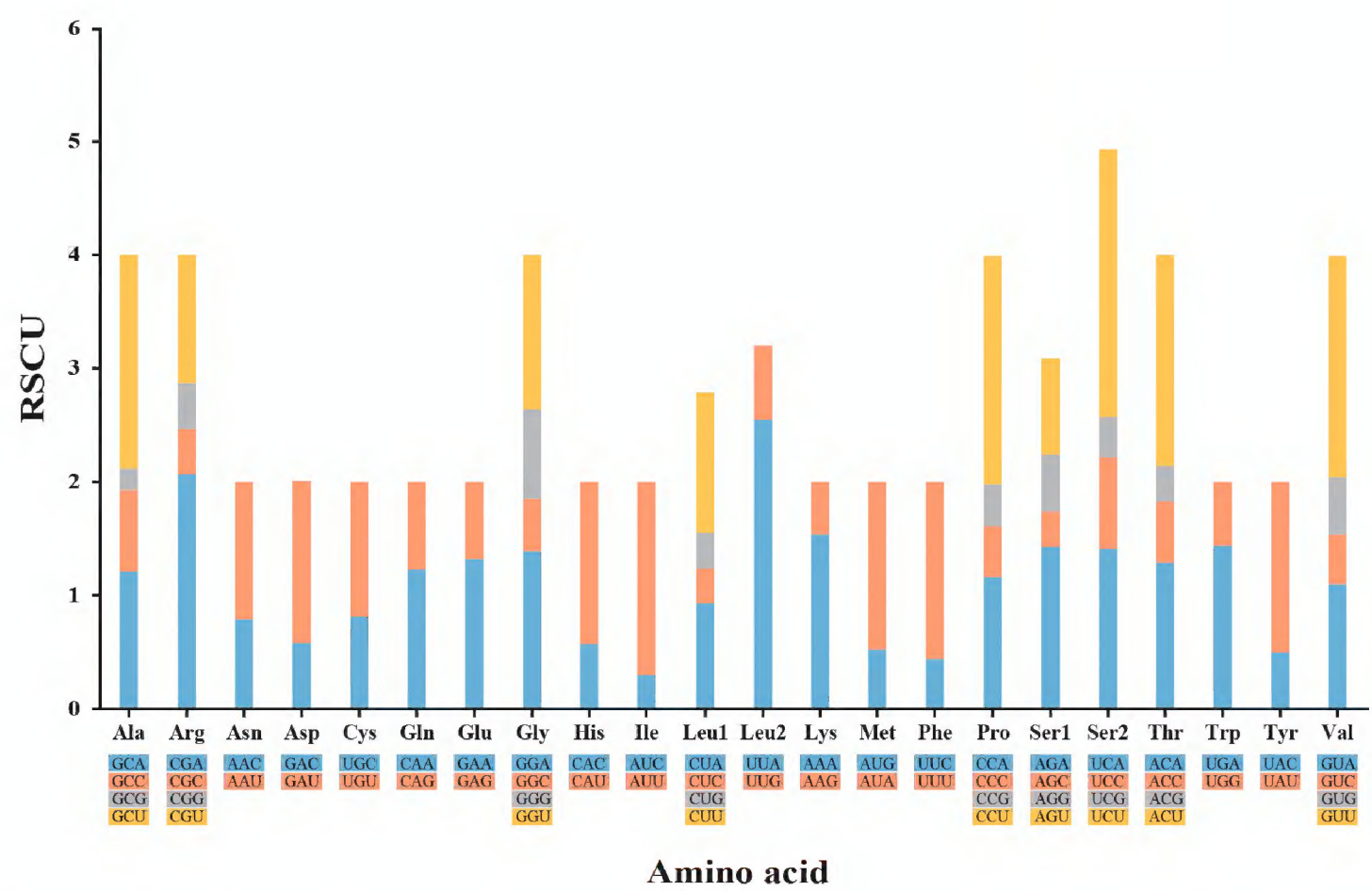
\*Termination codons were not included.



gene order of this genus is highly conserved (Bandyopadhyay et al. 2008; Cunha et al. 2009; Brauer et al. 2012; Barghi et al. 2016; Chen et al. 2016a, 2016b, 2016c; Gao et al. 2018; Uribe et al. 2018). The AT and GC-skew values of the entire genome sequences, which represent the measures of compositional asymmetry, were negative (−0.1233) and positive (0.0390), respectively, similar to those of cone snails (Gao et al. 2018).

PCGs and codon usage

The lengths of 13 PCGs of *C. lischkeanus* mitochondria range from 162 bp (*atp8*) to 1,716 bp (*nad5*) and contain 3,751 codons, excluding termination codons. The base composition of PCGs is 26.3% A, 39.2% T, 17.5% G, and 17.0% C, and the overall AT content was 65.5%, which is very similar to that of the entire mitochondrial genome sequence (AT content of 66.1%; Table 3). All PCGs have ATG as the initiation codon. With the exception of three PCGs (*nad1*, *nad4L*, and *nad4*) with TAG as a termination codon, all PCGs have TAA as a termination codon, which is consistent with complete mitochondrial genomes previously reported (Bandyopadhyay et al. 2008; Cunha et al. 2009; Brauer et al. 2012; Barghi et al. 2016; Chen et al. 2016a, 2016b, 2016c; Gao et al. 2018; Uribe et al. 2018). Fig. 2 shows the RSCU of *C. lischkeanus*, wherein the five most frequently used codons are UUA (Leu1), UCU (Ser2), CGA (Arg), CCU (Pro), and GUU (Val). In addition, codons with an A or U in the third position are the most frequently used, which is consistent with observations made in other mollusk species (Rawlings et al. 2010; Ren et al. 2010; Lee et al. 2019).



**Figure 2.** The relative synonymous codon usage (RSCU) frequency of the mitochondrial genome of *Conus lischkeanus*.



tRNA, rRNA genes, and D-loop regions

Twenty-two tRNA genes were found in the mitochondrial genome of *C. lischkeanus*. The length of tRNA genes range from 65 bp (*trnC*, *trnE*, *trnS2*, and *trnF*) to 70 bp (*trnL1*) (Table 2). All tRNA genes formed typical clover-leaf secondary structures, except for *trnS1* and *trnS2* which lack or had an imperfect D-arm (Fig. 3), which is common to other mollusk species (Boore 2006; Feng et al. 2020). Meanwhile, two ribosomal RNA genes with a total length of 2,327 bp consisting of small rRNA (*rrnS*; 952 bp) and large rRNA (*rrnL*; 1,375 bp) are located between *trnE* and *trnV*, and between *trnV* and *trnL1*, respectively (Fig. 2, Table 2). The D-loop is 587 bp in length and is located between *trnF* and *cox3*, with a short, inverted repeat (IR1; 20 bp), a typical feature of the mitochondrial genome of cone snail species. In contrast, the AT tandem repeat stretch found in *C. consors* G. B. Sowerby I, 1833 and *C. quercinus* [Lightfoot], 1786 was not identified in the *C. lischkeanus* mitochondrial genome (Brauer et al. 2012; Gao et al. 2018).

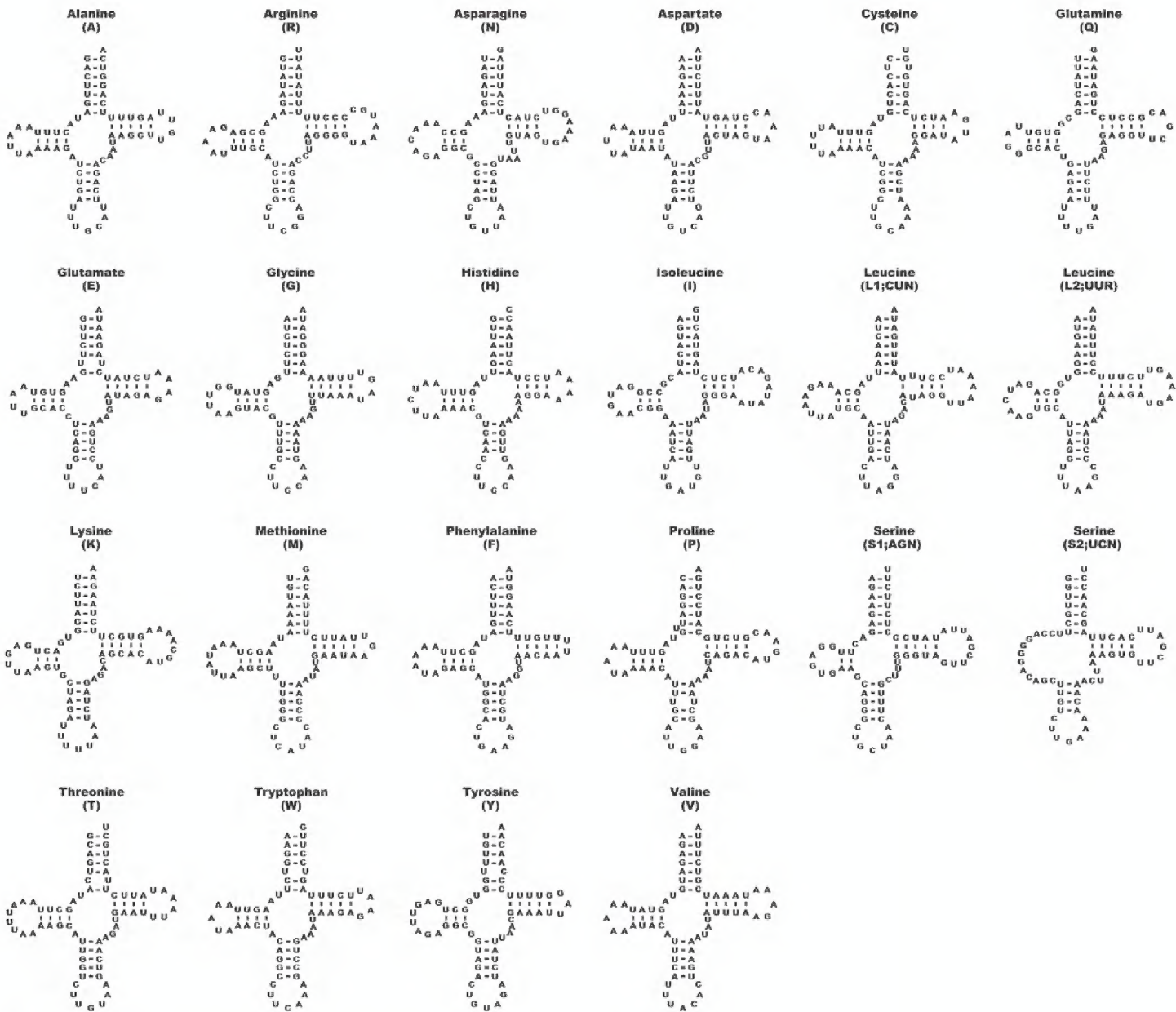
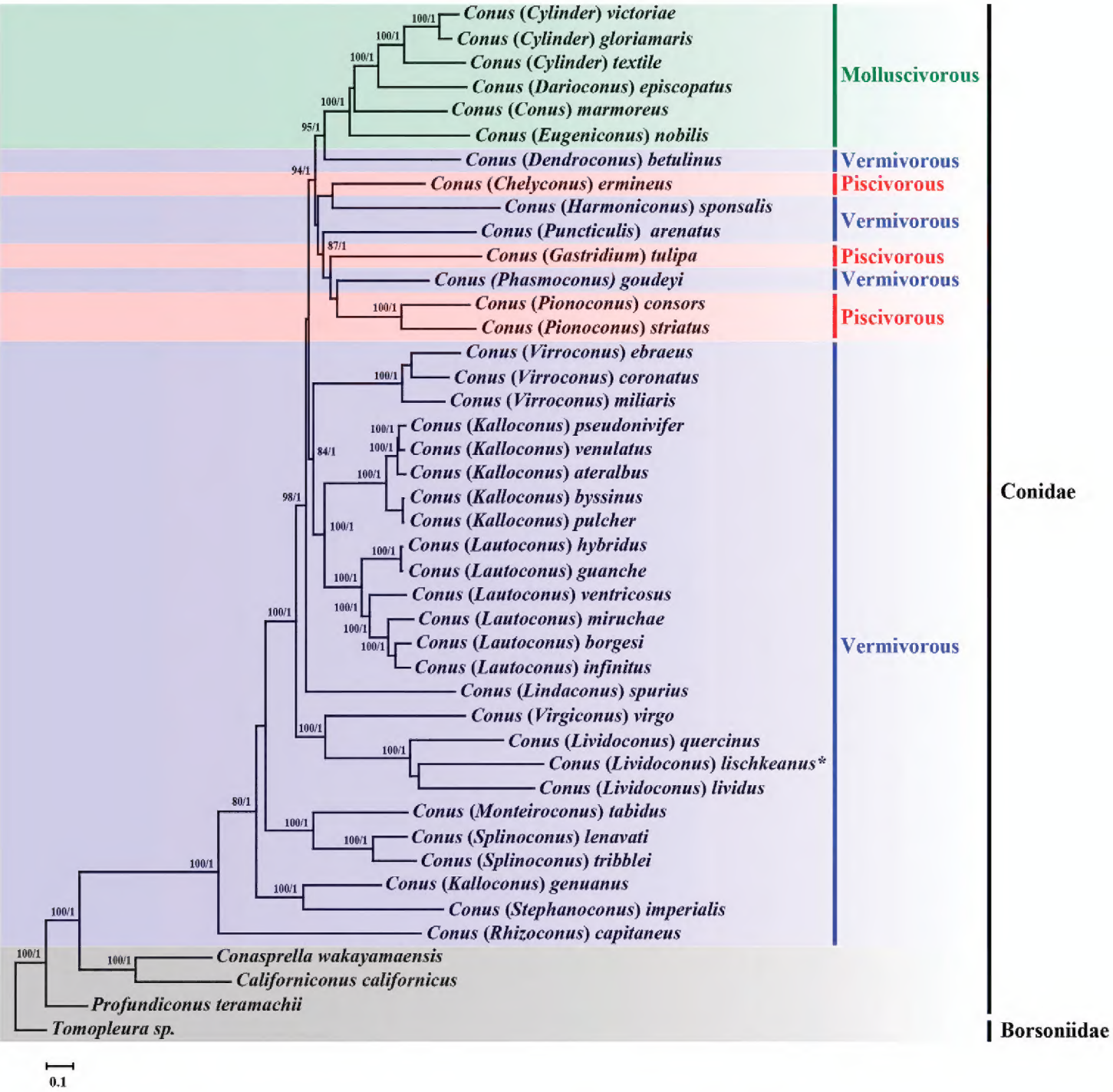


Figure 3. Predicted tRNA structures of *Conus lischkeanus*.



Phylogenetic implication of the evolutionary diversification of dietary specification

Phylogenetic analysis using ML and BI methods yield similar results with respect to the tree topology, as shown in Fig. 4. All subgenera, except *Kalloconus* da Motta, 1991, were monophyletic. A group of three *Conus* species, (*C. capitaneus*+(*C. imperialis*+*C. genuanus*)) was positioned at the most basal, but the branch reflected relatively weak supporting values (< 70% bootstrap values). Instead, the next monophyletic group consisting of (*C. tabidus*+(*C. lenavati*+*C. tribblei*)) was strongly supported (100% in ML and 1.0 BPP). Moreover, three species belonging to the subgenus *Lividoconus* Wils, 1970 (including *C. lischkeanus*) were grouped together with the subgenus *Virgiconus* Cotton,



**Figure 4.** Phylogenetic relationships of the genus *Conus* based on concatenated nucleotide sequences (13 protein coding genes plus two rRNA genes). Numbers above branches are statistical support values for ML (bootstrap values, > 70)/BI (posterior probability values, > 0.7). \*: determined in this study.



1945 species *C. virgo* Linnaeus, 1758, a sister to a large assemblage of the remaining *Conus* species that is composed of two well-supported groupings differing in their feeding type: vermivorous species and a mixture of three diet types. The “vermivorous only” clade is composed of three monophyletic groups of the subgenera *Virroconus* Iredale, 1930, *Kalloconus* da Motta, 1991, and *Lautoconus* Monterosato, 1923, with the latter two more closely related to each other than to *Virroconus*. Within the “mixed diet” clade, aside from a well-supported molluscivorous species (100% BP in ML and 1.0 BPP in BI), all vermivorous species are grouped either with piscivorous or vermivorous species. It is evident that vermivorous species are not monophyletic and are split into four branches, each forming sister relationships with other molluscivorous and/or piscivorous species. Given the mitochondrial genome phylogeny with vermivorous species positioned at the basal, the tree topology coincides with earlier hypothesis that worm-hunting was the ancestral diet type. Meanwhile, the other two diet types such as molluscivorous and piscivorous were secondarily derived (Duda Jr et al. 2001; Puillandre et al. 2014; Gao et al. 2018; Abalde et al. 2019). Notably, piscivorous species in our phylogenetic tree are not monophyletic and split into three branches, which is not consistent with previous mitochondrial genome phylogeny where fish-hunting species formed a monophyletic group (Gao et al. 2018). The polyphyly of piscivorous species in the current study implies that the fish-hunting species have evolved independently from worm-hunting groups multiple times. The complete mitochondrial genome information of the worm-hunting *Conus* species (*C. lischkeanus*) in the present study provides valuable insights into the mitochondrial genome diversity and molecular phylogeny of *Conus* species.

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